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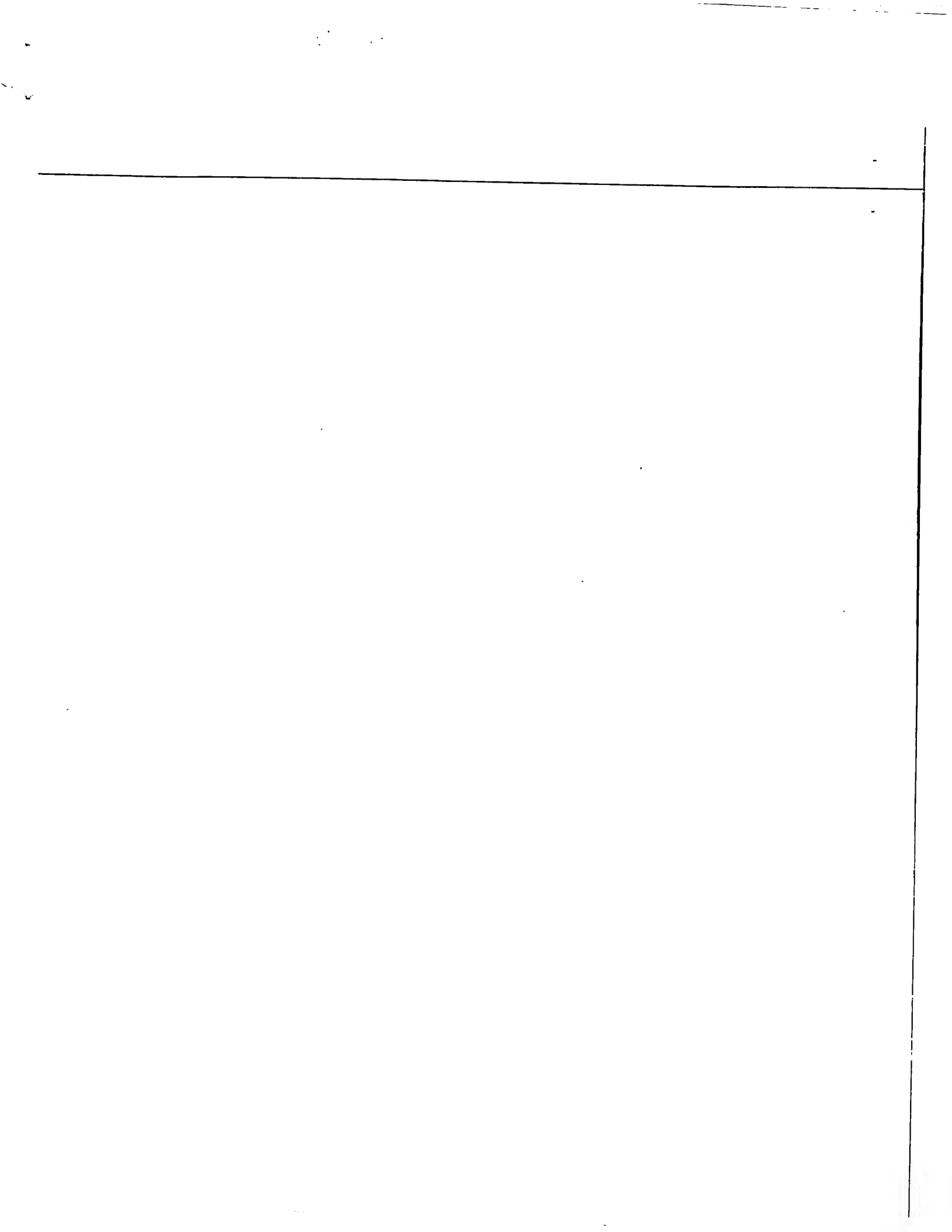
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Radiation-curable compositions for pigmented inks

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RADIATION-CURABLE COMPOSITIONS FOR PIGMENTED INKS

5 The present invention relates to radiation-curable compositions useful as ink vehicles or inks, in particular UV-curable inks, such as UV-litho inks.

UV-litho inks based on fatty acid modified polyester acrylate have been used since the 70ies with increasing success. Compared to conventional ink systems UV-inks exhibit
10 faster curing versus oxidative drying and provide an easier printing process because of no drying on the rollers, which leads to short start-up times and less waste. However, fatty acid modified polyester acrylate-based inks still show several performance shortcomings compared to conventional ink systems like the ink-water balance (lack of ink-water emulsion stability on press), pigment wetting (lack of flow in ink duct),
15 higher tack (picking) and higher misting, especially at higher press speeds.

A process for coating articles wherein a polymerizable prepolymer blend of an unsaturated ester of a vinyl aromatic hydrocarbon unsaturated alcohol copolymer such as styrene/allyl alcohol copolymer and a polymerizable interpolymmer ester such
20 as an unsaturated ester of a styrene/hydroxyalkyl acrylate/alkyl acrylate terpolymer is applied to the surface of the article to be coated and the prepolymer blend polymerized to form the coating using ionizing radiation is disclosed in US 3,546,002.

US 4,045,516 describes styrene-allyl alcohol copolymer-based solid polyene
25 compositions which when mixed with liquid polythiols can form solid curable polyene-polythiol systems. These solid polyenes, containing at least two reactive carbon-to-carbon unsaturated bonds, are urethane or ester derivatives of styrene-allyl alcohol copolymers.

30 French patent number 2.168.623 discloses styrene allyl alcohol copolymer-based urethane acrylates. The allylic hydroxyls are reacted with diisocyanate and capped with hydroxy acrylates.

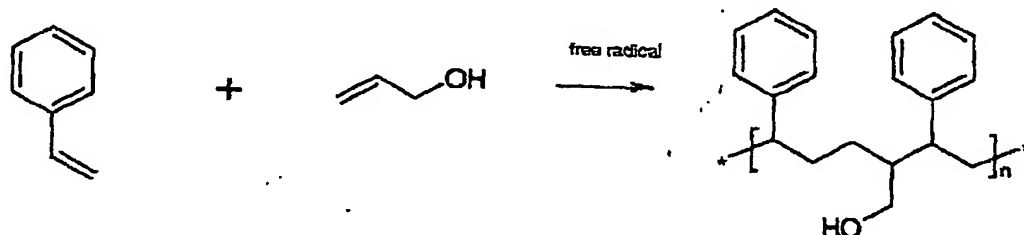
JP 49-94788 discloses the synthesis of styrene allyl alcohol (SAA) acrylates diluted in
35 trimethylolpropane trimethacrylate (TMPTA) as UV vehicles for printing inks.

While the known UV-litho inks exhibit good performance in some of their parameters, there is still a need for further ink vehicles and inks having improved properties such as an improved water balance, an improved pigment wetting and/or an improved cure speed.

The subject inventors have now surprisingly found that if acrylated SAA copolymer-based radiation-curable ink vehicles or inks are diluted with an alkoxyated acrylated monomer instead of an acrylated monomer such as TMPTA, water balance, pigment wetting and/or cure speed of the obtained ink vehicle or ink are improved.

Therefore, the present invention relates to a radiation-curable composition comprising an acrylated styrene allyl alcohol (SAA) copolymer and an alkoxyated acrylated monomer.

SAA copolymers are known in the art. They are for example produced by Lyondell via a radical polymerization of styrene and allyl alcohol as shown in the following scheme:



20

A process for making allylic/alkylenic copolymers is for example disclosed in WO 99/62969.

SAA copolymers are commercially available from Lyondell in three grades, SAA-100, SAA-101 and SAA-103. The properties of these copolymers which are preferred for the preparation of the present compositions are summarized in the following table 1.

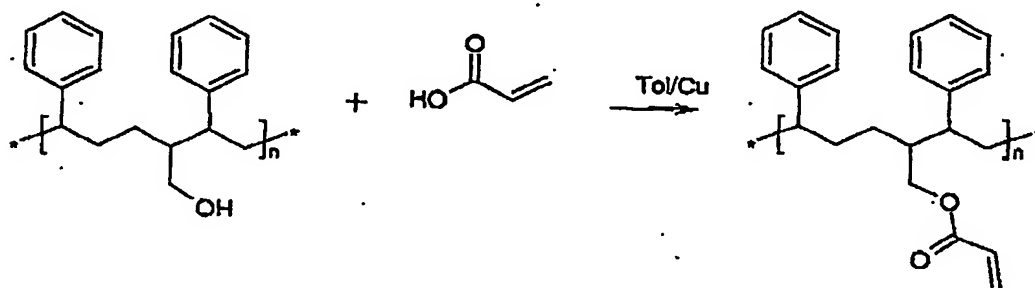
Table 1

Composition	SAA-100	SAA-101	SAA-103
Styrene mole %	70	60	80
Allyl alcohol mole %	30	40	20
Physical properties			
Physical form	Pastilles	Pastilles	Pastilles
color, Apha(30% in MEK)	40	40	40
Mn, GPC	1500	1200	3200
Mw, GPC	3000	2500	8400
Pd	2	2,1	2,6
OH number, mg KOH/g	210	255	125
Acid number, mg KOH/g	<0,1	<0,1	<0,1
Tg, °C	62	57	78
Softening point, °C	79	73	90

5

The hydroxyl groups of the SAA copolymers are stable to acids and bases and easily converted to acrylate by direct esterification with (meth) acrylic acid or by transesterification with an alkyl(meth)acrylate. The SAA copolymer can for example be dissolved in toluene and then reacted with acrylic acid or methacrylic acid under known process parameters as exemplified in the following scheme:

10



For the purpose of the present invention all acrylated copolymers and monomers may be derived from (meth)acrylic acid or from alkyl(meth)acrylates. For reasons of simplicity these compounds are uniformly called "acrylated", but it is to be understood that acrylated copolymers and monomers encompass acrylated and methacrylated copolymers and monomers throughout the present description and claims.

For reasons of processability and handling, SAA copolymer acrylates are preferably produced in the presence of a diluent. This diluent can be added during (polyol or acrylated

polyol) or after the acrylation step (only acrylated polyol). It has been found that alkoxyated acrylated monomers are particularly suitable as diluents for acrylated SAA copolymers. Moreover, litho inks based on acrylated SAA copolymers with alkoxyated acrylated monomers as diluents show exceptional performance compared to state of the art UV inks for water balance and pigment wetting. Additionally, depending on the alkoxyated acrylated monomer a composition exhibiting a very high cure speed can be obtained. Besides the SAA and the diluent, a third component such as an acidic component, preferably a fatty acid, can be added to modify the properties.

- 10 A monomer is herein defined as a component able to be further polymerized. It can be di or tri-functional but also tetra, penta or even hexafunctional.

Generally, the composition of the present invention should comprise the minimum amount of 10% by weight, more preferably a minimum amount of 40% by weight and most preferably a minimum amount of 50% by weight alkoxyated acrylated monomer, based on the total weight of the acrylated SAA copolymer and the monomer. For example the composition may comprise 10-90% by weight acrylated SAA copolymer and 10-90% by weight alkoxyated acrylated monomer, preferably 40-70% by weight acrylated SAA copolymer and 30-60% by weight alkoxyated acrylated monomer, each based on the total weight of copolymer and monomer.

The acrylic or methacrylic monomer derivatives used as diluents in the compositions of the present invention are those currently used in radiation-curable compositions, especially in the ink field. They can be chosen among mono-, di-, tri- and polyhydroxy compounds being alkoxyated and acrylated. The preferred alkoxyates are ethoxyates and propoxyates. Each of the hydroxy groups may bear one or more, such as two or three, alkoxy moieties being acrylated or methacrylated.

Advantageously, the alkoxyated acrylated monomer used in the composition of the present invention is selected from the group consisting of alkoxyated triol triacrylates, alkoxyated diol diacrylates and alkoxyated acrylated bisphenol A derivatives, such as ethoxyated glycerol triacrylates, propoxyated glycerol triacrylates, ethoxyated neopenthyglycol diacrylates, propoxyated neopenthyglycol diacrylates, ethoxyated trimethylolpropane triacrylates, propoxyated trimethylolpropane triacrylates, acrylated bisphenol A ethoxyates, and acrylated bisphenol A propoxyates. Particularly preferred monomer diluents are acrylated bisphenol A alkoxyates such as acrylated bisphenol A ethoxyate, available as EbecrylTM 150 from Surface Specialties-UCB. Propoxyated glycerol triacrylate is for example available under the trademark OTA480.

The radiation-curable composition of the present invention is useful as ink vehicle for the preparation of inks, such as UV-litho inks. Therefore, the composition may further comprise one or more compounds selected from pigments, photoinitiators, and ink additives.

The pigments used in the compositions of the invention are every pigments used in paste or liquid inks. A list of such pigments can be found in the Color Index. More particularly, those pigments may be cited such as Process Yellow 13 (Diarylide Yellow - Irgalite BAW of Ciba, Permanent GR. of Clariant), Process Magenta Pigment 57 (Bona Calcium - Ilobona 4BY of Sun, Irgalite SMA of Ciba), Process Blue 15.3 (Copper Phthalocyanine - Irgalite GLO of Ciba, Hostaperm Blue B2G of Clariant), Process Black 7 (Oxidised Carbon Black - Special Black 250; Special Black 350 of Degussa), etc. The pigments are preferably used at 1-50% by weight of the total weight of the composition, more preferably at 1-40 % by weight.

The photoinitiators usable in the compositions of the invention are well known in the art. They can be chosen from α -hydroxyketones, α -aminoketones, benzildimethyl-ketals, acyl phosphines, benzophenone derivatives, thioxanthenes and blends of these. They are used at 0 to 15% by weight. Photoactivators are chosen between amine derivatives. The photoinitiators need only be used if the compositions are cured by ultraviolet light. The compositions may also be cured by electron beams rays, and, in this case, no photoinitiator needs to be added to the composition.

The additives are those commonly used in inks, such as stabilizers, substrate wetting agents, anti-foam agents, adhesion promoters, dispersing agents etc.

The compositions of the present invention may comprise each one acrylated SAA copolymer and one alkoxyated acrylated monomer or, if desired, two or more different acrylated SAA copolymers and two or more different alkoxyated acrylated monomers.

The compositions of the present invention can be prepared by acrylating SAA copolymer as described above. Work-up of the reaction product is, however, difficult due to the high viscosity of the acrylate. To ensure efficient washes and phasing of the acrylated SAA copolymer, a diluent in the form of the alkoxyated acrylated monomer can be added directly after acrylation, but before the solvent removal (acrylation of SAA with post dilution with (meth)acrylated polyols or "post dilution process").

In a variant, the SAA can be acrylated in presence of (meth)acrylated polyols.

It has now surprisingly been found that if the SAA copolymer and the diluent in the form of an alkoxyated monomer are acrylated simultaneously in-situ (co-acrylation process) better properties can be obtained such as ink water balance. The positive impact on monomer hydroxyl conversion of the co-acrylation process compared to the post dilution process is demonstrated by the LCMS data summarized in the following table 2:

Resin	Process	SAA	Diluent		Monomer composition (LC area%)		
			Type	%	Mono	Di	Tri
1	Co-acrylation	103	OTA480	50	0.8	5.4	93.8
2	Post-dilution	103	OTA480	50	1.2	10.4	88.4
3	Co-acrylation	100	Eb150	60	7	93	NA
4	Post-dilution	100	Eb150	60	14	86	NA
5	Co-acrylation	103	Eb150	70	5	95	NA
6	Post-dilution	103	Eb150	70	12	88	NA

- 10 The above table shows that bisphenol A ethoxylate (Ebecryl™ 150) is less monoacrylated and propoxylated glycerol (OTA480) is less mono- and diacrylated (based on LC% area) in the co-acrylation process compared to the post dilution process. While applicant does not wish to be bound to any theory, it is believed that this effect can probably explain the improvement in the ink-water balance of compositions prepared according to the co-acrylation process compared to the compositions prepared according to the post dilution process.

Remarkably, there is also a difference in appearance when the co-acrylation process is used. The obtained resin is much less colored.

20 Furthermore, the cure speed of the compositions of the present invention has been measured in a cyan ink with 8% photoinitiator blend (32% EDB/13% DETX/7% Irgacure369/14% benzophenone/34%BDK) under a 120 Wcm Hg lamp by graphite test. The results are summarized in the following table 2a:

SAA	Process	Diluent	Alkoxylation	Cure speed m/min
100	co-acrylation	EB150:65%	OE	100
100	post dilution	EB150:60%	OE	90
100	co-acrylation	EB150:60%	OE	90
100	co-acrylation	OTA480:45%	OP	40
100	co-acrylation	BisA(OP)2DA	OP	70
100	co-acrylation	BisA(OP)4DA	OP	60
100	co-acrylation	TMP(EO)3TA	OE	25
103	co-acrylation	EB150:70%	OE	110
103	post dilution	EB150:70%	OE	90
103	post dilution	OTA480:50%	OP	20

It can be seen from the above table 2a that the cure speed is not influenced by the process used (co-acrylation or post dilution). Surprisingly, cure speed of Bisphenol A monomers is greater than for other monomers such as OTA 480 or alkoxyated TMPTA. Among the Bisphenol A monomers, ethoxylated Bisphenol A monomers have a higher cure speed and are thus preferred over propoxylated Bisphenol A monomers.

Inks are generally made in 2 steps, the pigment dispersion step and the letdown step. In the 1st step, all the pigments are added in the pigment dispersing binders (oligomers and monomers). They are mixed and then dispersed on a triple roll or bead mill. A few passes are necessary to achieve a good dispersion. Pigments that are difficult to disperse require more number of passes. It is the same with the binders. Binders that are not good for pigment wetting also require additional passes. Once the pigment has achieved this fineness, the pigment paste is diluted with the letdown. The letdown has to be compatible with the binder used to disperse the pigments. The finished ink is then printed onto the substrate. The ink film is then cured under a UV lamp, for example at 120W/cm and 30 m/min. A few passes may be required to cure the ink if the binder is not reactive enough.

The invention also relates to the polymeric compositions obtainable by curing the radiation curable composition as well as to substrates being partially or entirely coated with the polymeric composition.

Pigment wetting can be evaluated by different methods:

- Rheology: Pigment wetting is a major factor of influence on the rheology. Inks with bad wetting of the pigment are showing a marked shear thinning effect, whereby the viscosity is high at low shear rate and drops as the shear rate is increased. This results in a high shortness index (SI = ratio of low shear viscosity to high shear viscosity). For liquid inks a Newtonian rheology is required. Ideally, this means that the viscosity is independent of the shear rate. (SI = 1). Paste inks are more pseudoplastic, showing a shear depending viscosity. (SI > 1). Too high SI (too high low shear viscosity) may result in bad flow in the ink duct. The rheology is measured with cone and plate type rheometers.
- Optical density: Pigment wetting can also be evaluated by measuring the color density of the printed ink at constant film thickness. In this case the ink is printed using a lab applicator and the color density is measured with a densitometer, which spectrophotometrically compares the reflected light to the incident light.
- For the present invention the pigment wetting is rated on a scale from 5 = excellent to 0 = bad pigment wetting.
- The water balance of the compositions of the present invention was evaluated on lithotronic.
- Basically, the Lithotronic measures the torque needed for a certain speed (rpm). The torque gives a measure for viscosity. With the Lithotronic, the change in viscosity of an ink is measured when water is emulsified in it.
- The measurement consists of two phases: preconditioning and measurement. During preconditioning, the sample is sheared at constant speed and heated at the same time to a certain preprogrammed temperature. At the end of the preconditioning phase, the sample has reached a stable viscosity. At that moment, controlled metering of fountain solution is started. Changes of applied torque (hence viscosity) versus time and emulsion capacity are recorded. When maximum emulsion capacity is reached, a drop in torque is usually experienced because of the free water in the beaker.
- At first contact with water, change of torque (ΔT) should be small. Further, when water is emulsified in the ink, viscosity should only undergo a minor increase. This ensures a good ink transfer on the press. If the emulsion is too fine and too stable (too high increase of viscosity), it will lead to a loss of density and possible ink build up. If the emulsion is too coarse (viscosity decrease), it can lead to unstable press behaviour making regular press control necessary.
- For the present invention the ink water balance is rated by the type of emulsion (F = good ink water balance characterized by a limited viscosity increase, resulting from a fine

emulsion; C = bad ink water balance characterized by a high viscosity decrease, resulting from a coarse emulsion).

5 The invention will now be illustrated by the following non-limiting examples and tests which are by way of illustration only. Unless otherwise indicated, herein all the test results and properties herein were performed using conventional methods well known to those skilled in the art. The amounts in the tables are given in % by weight based on the total weight of the composition.

10 The compositions of the present invention were prepared according to the following production examples:

Composition 1

15 Charge a reactor with 524 grams of SAA-100 from Lyondell (1,96 eq), 921,4 grams Simulsol BPJE (ethoxylated bisphenol A from Seppic - Air Liquide) (4,60 eq), 496,4 grams acrylic acid (6,89 eq), 1295 grams toluene, 48,5 grams para-toluenesulphonic acid, 0,50 grams Cu₂O, 9,7 grams hypophosphorous acid. Heat the reactor content at 120°C until the rate
20 of reaction slows down as measured by collection of the esterification water. After about 8 to 10 hrs when the acid value is less than 15, cool down to 50°C and dilute with toluene till a 1/1 reactant:solvent ratio. Charge the reactor with an aqueous solution of 15% sodium chloride and stir for 2 minutes. Let phase separate at 40-50°C and decant the aqueous
25 phase. Repeat saline washes until an acid value of 5 is reached. Charge the reactor with a caustic solution of 5% NaOH in an amount of 25% of the batch weight and stir for 2 minutes. Phase separate at 40°C and decant the aqueous layer. Then add 1,8 grams HQ and 3,6 grams trisnonylphenylphosphite and heat till 120°C to distill the remaining water. Maintain an airsparge of 1 l/hr. When the water content is below 0,1% cool down for
30 filtration. After distillation of the toluene at 90°C and a reduced pressure of 30 mbar, a product A1 with a Cone & Plate viscosity of 33 Pas and an acid value of 0,7 mg KOH/g is obtained.

Composition 2

35 Charge a reactor with 399,9 grams of SAA-103 from Lyondell (0,90 eq), 65,0 grams acrylic acid (0,90 eq), 313 grams toluene, 11,8 grams para-toluenesulphonic acid, 0,07 grams Cu₂O, 2,35 grams hypophosphorous acid. Heat the reactor content at 120°C until the rate

- of reaction slows down as measured by collection of the esterification water. After about 8 to 10 hrs when the acid value is less than 15, cool down to 50 °C and dilute with toluene till a 1/1 reactant:solvent ratio. Charge the reactor with an aqueous solution of 15% sodium chloride and stir for 2 minutes. Let phase separate at 40-50°C and decant the aqueous phase. Repeat saline washes until an acid value of 5 is reached. Charge the reactor with a caustic solution of 5% NaOH in an amount of 25% of the batch weight and stir for 2 minutes. Phase separate at 40°C and decant the aqueous layer. Then, add 1050 grams acrylated bisphenol ethoxylate (EB 150 from Surface Specialties - UCB), 1,5 grams HQ and 3,0 grams trisnonylphenylphosphite and heat till 120°C to distill the remaining water. Maintain an airsparge of 1 l/hr. When the water content is below 0,1% cool down for filtration. After distillation of the toluene at 90°C and a reduced pressure of 30 mbar, a product B1 with a Cone & Plate viscosity of 53 Pas and an acid value of 1.05 mg KOH/g is obtained.
- Further compositions were prepared according to the above production examples. The following compositions (products A1-A5 and B1-B2) were obtained according to the invention:

Table 3

Example	SAA copolymer	Diluent		Viscosity Pas; 25°C C&P	Acid value Mg KOH/g
		Type	level		
Product A1	SAA 100	Bis A (EO)4 DA	65%	33	0,7
Product A2	SAA 100	Bis A (FO)2 DA	80%	75	0,8
Product A3	SAA 100	Bis A (PO)4 DA	75%	130	0,8
Product A4	SAA 100	TMP (EO)3 TA	45%	40	1,2
Product A5	SAA 100	Glycerol (PO)3 TA	45%	87	0,8
Product B1	SAA 103	Bis A (EO)4 DA	70%	53	1,05
Product B2	SAA 103	Glycerol (PO)3 TA	50%	117	0,8

In the above table 3 the following abbreviations are used:

- Bis A: bisphenol A
 TMP: trimethylolpropane
 EO: ethyleneoxide
 PO: propyleneoxide
 DA: diacrylate

TA: triacrylate

C&P: Cone and Plate

The following comparative compositions were prepared as well:

5 Table 4

Example	Polymer	Diluent		Viscosity Pas; 25°C C&P	Acid value Mg KOH/g
		Type	level		
Product C1	SAA 100	TMPTA	50%	33	0.9
Product C2	SAA 100	HDDA	30%	75	0.9
				130	
Product C3	PEA:Eb657	NA	NA	40	<15
Product C4	PEA:Eb811	NA	NA	87	<1

In the above table 4 the following abbreviations are used:

- 10 PEA:Eb657: Polyester acrylate Ebecryl™ 657
 PEA:Eb811: Polyester acrylate Ebecryl™ 811
 TMPTA: trimethylolpropane trimethacrylate
 HDDA: Hexane diol diacrylate
 NA: not applicable.

15

With the above compositions and comparative compositions the following ink formulations were prepared:

Table 5

	A1	A2	A3	A4	A5	B1	B2	C1	C2	C3	C4
Oligomer	64	50	62	59	60	58	58	64	66	57	60
Stab	1	1	1	1	1	1	1	1	1	1	1
PB 15:3	17	17	17	17	17	17	17	17	17	17	17
Talc	6	6	6	6	6	6	6	6	6	6	6
PI blend	8	8	8	8	8	8	8	8	8	8	8
OTA480					8		10			11	8
TMPTA								4			
HDDA									2		
TMPE(O)3TA				9							
EbecrylTM 150	4					10					
BPA(OP)2DA		18									
BPA(OP)4DA			6								

Stab: polymerisation inhibitor (5% solution of NPAL in DPGDA (DiPropyleneGlycolDiAcrylate); NPAL = Tris(N-nitroso-N-phenylhydroxylamine)aluminium salt)

PB15:3 = phthalocyanine pigment (IrgaliteTM GLO from Ciba)

Various properties of the obtained ink formulations were measured. The results are summarized in the following table 6:

Table 6

	A1	A2	A3	A4	A5	B1	B2	C1	C2	C3	C4
Visco 2.5 s-1	62.2	61.5	63.0	75.0	73.1	58.1	76.8	71.2	131	72.0	71.0
Visco 100 s-1	35.1	34.8	33.1	34.5	34	32.5	34.3	30.4	37.6	30.0	30.2
SI	1.8	1.8	1.9	2.2	2.1	1.8	2.2	2.3	3.5	2.4	2.4
Optical Density	2.05	1.99	2.08	1.98	2.08	2.08	2.13	2.1	1.7	1.95	2.04
Pigment wetting	4	4	4	3	3	4	3	2	1	2	2
Delta T	9	8	8	9	12	9	8	9	5	23	20
Emulsion	F	F	F	F	F	F	F	C	C	C	C
Cure speed	100	70	60	25	40	110	30	40	50	20	20

Cure speed: m/min at 120W/cm (graphite test)

Furthermore, a comparison between product A1 of the present invention and an ink formulation based on polyesteracrylate (Ebecryl657) was conducted with various pigments. The ink formulations prepared are summarized in the following table 7:

Table 7

	Black		Cyan		Magenta		Yellow	
	A1	Eb657	A1	Eb657	A1	Eb657	A1	Eb657
Oligomer	63.5	59	64	57	64	59	68	64
Stab	1	1	1	1	1	1	1	1
P.Black 7	20	20						
P.Blue 15:3			17	17				
P.Red 57:1								
P.Yellow 13					18	18		
P.Blue 61:1	1.5	1.5					14	14
Talc			6	6	3	3		
PI blend	8	8	8	8	8	8	8	8
OTA480		10.5		11		11		
Eb150	6		4		6		6	10

5 Various parameters of the obtained inks were measured. The results are summarized in the following table 8:

Table 8

	Black		Cyan		Magenta		Yellow	
	A1	Eb657	A1	Eb657	A1	Eb657	A1	Eb657
Visc. 2.5 s-1	55.9	102	62.2	72.0	40.3	60.1	97.4	87
Visc. 100 s-1	32.1	31.7	35.1	30.0	33.9	33.4	34.8	34.8
SI	1.7	3.2	1.8	2.4	1.2	1.8	2.8	2.5
Optical density	2.35	1.59	2.05	1.95	1.8	1.84	1.65	1.60
Pigment wetting	4	0	4	2	5	3	2	2
Delta T (%)	9	19	9	23	6	10	7	20
Emulsion	F	C	F	C	F/C	C	F	C
Cure speed	70	30	100	20	80	30	70	20

10 Cure speed: m/min at 120W/cm (graphite test)

15 The above results demonstrate that the compositions of the present invention exhibit an improved water balance and an improved pigment wetting compared to acrylated SAA copolymer-based ink vehicles which do not comprise an alkoxyated acrylated monomer and an improved water balance and an improved pigment wetting compared to polyester acrylate ink vehicles. Moreover, the cure speed of the radiation-curable composition according to the present invention can be higher compared to the known compositions depending on the alkoxyated acrylated monomer employed.

Claims

1. Radiation-curable composition, comprising an acrylated styrene allyl alcohol (SAA)
5 copolymer and an alkoxyated acrylated monomer.
2. Composition according to claim 1, comprising 10-90% by weight acrylated SAA
copolymer and 10-90% by weight alkoxyated acrylated monomer, each based on the total
10 weight of the acrylated SAA copolymer and alkoxyated acrylated monomer.
3. Composition according to any of the preceding claims, wherein the alkoxyated
acrylated monomer is selected from the group consisting of alkoxyated triol triacrylates,
alkoxyated diol diacrylates, preferably alkoxyated acrylated bisphenol A derivatives, more
15 preferably oxyethylated.
4. Composition according to claim 3, wherein the alkoxyated acrylated monomer is
selected from the group consisting of ethoxylated glycerol triacrylates, propoxylated glycerol
triacrylates, ethoxylated neopenthyglycol diacrylates, propoxylated neopenthyglycol
diacrylates, ethoxylated trimethylolpropane triacrylates, propoxylated trimethylolpropane
20 triacrylates, acrylated bisphenol A ethoxylates, acrylated bisphenol A propoxylates,
alkoxyated ditrimethylolpropane tetraacrylates, alkoxyated pentaerythritol tetraacrylates
and alkoxyated dipentaerythritol penta/hexa acrylates.
5. Composition according to any of the preceding claims, further comprising one or
25 more compounds selected from pigments, photoinitiators, and ink additives, such as
stabilizers, substrate wetting agents, anti-foam agents, adhesion promoters, dispersing
agents etc.
6. Process of preparing a radiation-curable composition, comprising the steps of
30
 - a) admixing a styrene allyl alcohol (SAA) copolymer with an alkoxyated polyol
and
 - b) in-situ acrylating the mixture obtained under a).
7. Radiation-curable composition obtainable by the process of claim 6.
35
8. Use of the radiation-curable composition as claimed in any of claims 1-5 or 7 as ink
vehicle or ink.

9. Polymeric composition obtainable by curing the radiation-curable composition as claimed in any of claims 1-5 or 7.
10. Substrate partially or entirely coated with the polymeric composition as claimed in claim 9.

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Abstract

The present invention relates to radiation-curable compositions useful as ink vehicles or inks, in particular UV-curable inks, such as UV-litho inks.

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